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**INFORMATION DISCLOSURE
STATEMENT BY APPLICANT**

(Use as many sheets as necessary)

Complete if Known

Application Number	10/687,677
Filing Date	10/17/2003
First Named Inventor	GUY
Art Unit	1042 1632
Examiner Name	/Wu Cheng Winston Shen/ (12/04/2006)
Attorney Docket Number	5853-324

Sheet	1	of	2
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OTHER PRIOR ART—NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
W.S.		Larsson NG, Andersen O, Holme E et al. Leber's hereditary optic neuropathy and complex I deficiency in muscle. Ann Neurol. 1991; 30:701-708	
W.S.		Majander A, Huoponen K, Savontaus ML et al. Electron transfer properties of NADH:ubiquinone reductase in the ND1/3460 and the ND4/11778 mutations of the Leber hereditary optic neuroretinopathy (LHON). FEBS Lett. 1991; 292:289-292	
W.S.		Vergani L, Martinuzzi A, Carelli V et al. MtDNA mutations associated with Leber's hereditary optic neuropathy: studies on cytoplasmic hybrid (cybrid) cells. Biochem Biophys Res Commun. 1995; 210:880-888	
W.S.		Wallace DC. Mitochondrial diseases in man and mouse. Science. 1999; 283:1482-1488	
W.S.		Carelli V, Ghelli A, Bucchi L et al. Biochemical features of mtDNA 14484 (ND6/M64V) point mutation associated with Leber's hereditary optic neuropathy. Ann Neurol. 1999; 45:320-328	
W.S.		Chinnery PF, Johnson MA, Wardell TM et al. The epidemiology of pathogenic mitochondrial DNA mutations. Ann Neurol. 2000; 48:188-193	
W.S.		Guy J, Qi X, Hauswirth WW. Adeno-associated viral-mediated catalase expression suppresses optic neuritis in experimental allergic encephalomyelitis. Proc Natl Acad Sci U S A. 1998; 95:13847-13852	
W.S.		Wallace DC, Singh G, Lott MT et al. Mitochondrial DNA mutation associated with Leber's hereditary optic neuropathy. Science. 1988; 242:1427-1430	
W.S.		Hartl FU, Neupert W. Protein sorting to mitochondria: evolutionary conservations of folding and assembly. Science. 1990; 247:930-938	
W.S.		Schon EA. Mitochondrial genetics and disease. Trends Biochem Sci. 2000; 25:555-560	

Examiner Signature	/Wu Cheng Winston Shen/ (12/04/2006)	Date Considered	
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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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W.S.		Guy J, Qi X, Muzyczka N et al. Reporter expression persists 1 year after adeno-associated virus- mediated gene transfer to the optic nerve. Arch Ophthalmol. 1999; 117:929-937	
W.S.		Esposito LA, Melov S, Panov A et al. Mitochondrial disease in mouse results in increased oxidative stress. Proc Natl Acad Sci U S A. 1999; 96:4820-4825	
W.S.		Brown MD, Trounce LA, Jun AS et al. Functional analysis of lymphoblast and cybrid mitochondria containing the 3460, 11778, or 14484 Leber's hereditary optic neuropathy mitochondrial DNA mutation. J Biol Chem. 2000; 275:39831-39836	
W.S.		Brown MD. The enigmatic relationship between mitochondrial dysfunction and Leber's hereditary optic neuropathy. J Neurol Sci. 1999; 165:1-5	
W.S.		Cock HR, Cooper JM, Schapira AH. Functional consequences of the 3460-bp mitochondrial DNA mutation associated with Leber's hereditary optic neuropathy. J Neurol Sci. 1999; 165:10-17	
W.S.		Sazanov, L. Resolution of the Membrane Domain of Bovine Complex I into Subcomplexes: Implications for the Structural Organization of the Enzyme. Biochemistry 2000, 39: 7229-7235	

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